

chain nodes :

30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 14 16 17 18 21 22 23 24 25 26

chain bonds :

2-41 2-46 3-42 3-45 4-9 8-33 10-32 11-31 12-30 14-43 14-44 17-39 17-40
21-37 22-38 24-34 25-35 26-36

ring bonds :

1-2 1-7 1-14 2-3 3-4 4-5 4-14 5-6 6-7 8-9 8-13 9-10 10-11 11-12 12-13
16-17 16-18 17-18 21-22 21-26 22-23 23-24 24-25 25-26

exact/norm bonds :

1-2 1-7 1-14 4-5 4-14 5-6 6-7 12-30

exact bonds :

2-3 2-41 2-46 3-4 3-42 3-45 4-9 8-33 10-32 11-31 14-43 14-44 16-17
16-18 17-18 17-39 17-40 21-37 22-38 24-34 25-35 26-36

normalized bonds :

8-9 8-13 9-10 10-11 11-12 12-13 21-22 21-26 22-23 23-24 24-25 25-26

isolated ring systems :

containing 1 : 8 : 16 : 21 :

G1:X, [*1], [*2]

Match level :

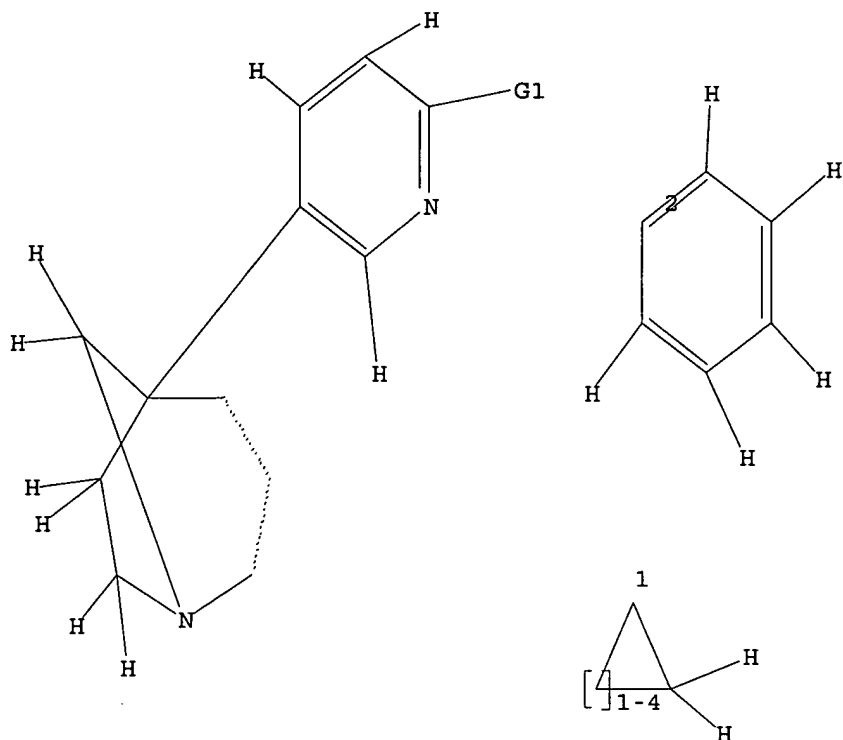
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34:CLASS 35:CLASS 36:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS
42:CLASS 43:CLASS 44:CLASS 45:CLASS 46:CLASS

L1 STRUCTURE UPLOADED

=> d

L1 HAS NO ANSWERS

L1 STR



G1 X, [@1], [@2]

Structure attributes must be viewed using STN Express query preparation.

=> s l1 ful
 FULL SEARCH INITIATED 12:35:59 FILE 'REGISTRY'
 FULL SCREEN SEARCH COMPLETED - 70 TO ITERATE

100.0% PROCESSED 70 ITERATIONS 10 ANSWERS
 SEARCH TIME: 00.00.01

L2 10 SEA SSS FUL L1

=> fil caplus
 COST IN U.S. DOLLARS SINCE FILE TOTAL
 ENTRY SESSION
 FULL ESTIMATED COST 166.94 167.15

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=> s l2
L3 1 L2

=> d bib abs

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2003:534333 CAPLUS

DN 139:101039

TI Derivatives of 5-(pyridin-3-yl)-1-azabicyclo[3.2.1]octane, their preparation, and their application in therapy as nicotinic receptor ligands for treatment of CNS disorders

IN Galli, Frederic; Leclerc, Odile; Lockheed, Alistair

PA Sanofi-Synthelabo, Fr.

SO Fr. Demande, 20 pp.

CODEN: FRXXBL

DT Patent

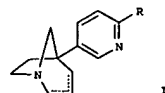
LA French

FAN.CNT 1

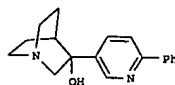
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PI FR 2834511	A1	20030711	FR 2002-109	20020107
FR 2834511	B1	20040213		
CA 2471628	AA	20030717	CA 2003-2471628	20030103
WO 2003057697	A1	20030717	WO 2003-FR4	20030103
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RW:	GH, GM, KE, LS, MM, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LJ, MC, ML, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2003216777	A1	20030724	AU 2003-216777	20030103
EP 1465893	A1	20041013	EP 2003-712202	20030103
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BR 2003006707	A	20050209	BR 2003-6707	20030103
JP 2005514422	T2	20050519	JP 2003-558012	20030103
US 2005020568	A1	20050127	US 2004-500015	20040623
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WO 2003-FR4	W	20030103		
OS MARPAT 139:101039				
GI				

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

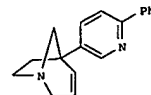
(Continued)



I



II



III

AB Title compds. I and their acid addition salts are disclosed [wherein: R = halo, a Ph group (substituted by one or more groups chosen from halo, Cl-6

alkyl or alkoxy, NO₂, amino, CF₃, cyano, OH, acetyl, or methylenedioxy), pyridinyl, thienyl, indolyl, or pyrimidinyl (possibly substituted by one or more Cl-6 alkoxy); dashed bonds = one single bond and another single

or double bond]. The compds. are useful as pharmaceuticals, particularly as CNS agents, and specifically as ligands of nicotinic receptors. The compds. were tested against nicotinic receptors with the α₄β₂ subunit, or with the α₇ subunit. Four synthetic examples and a list of 35 specific compds. (as either di- or tri-HBr or 1:1 oxalate salts)

are given. For instance, 2,5-dibromopyridine was arylated in the 2-position by PhB(OH)₂ using Pd(PPh₃)₄ catalyst, and the resultant 5-bromo-2-phenylpyridine was lithiated with BuLi and treated with 1-azabicyclo[2.2.2]octan-3-one to give the bicyclic alc. II. Dehydration and rearrangement of II by heating with MeSO₃H at 180° gave invention compound III, isolated as the di-HBr salt. In tests for specific

binding to isolated rat cerebral nicotinic receptors having either α₄β₂ or α₇ subunits, compds. I had IC₅₀ values in the ranges of 0.01-10 μM and 0.005-20 μM, resp. Some compds. showed selectivity for the α₇ receptor subtype.

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